



Sheet 1 of 1

SUBSTITUTE FORM PTO-1449 (MODIFIED)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		Attorney Docket No.	07540/020003	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Use several sheets if necessary)				Serial No.	09/292,862	
				Applicant	Michael A. Walter et al.	
				Filing Date	April 16, 1999	
				Group	1614	
(37 CFR §1.98(b))				IDS Filed	October 22, 1999	
U.S. PATENTS						
Examiner's Initials	Patent Number	Issue Date	Patentee	Class	Subclass	Filing Date (if Appropriate)
FOREIGN PATENT OR PUBLISHED FOREIGN PATENT APPLICATION						
Examiner's Initials	Document Number	Publication Date	Country or Patent Office	Class	Subclass	Translation (Yes/No)
OTHER DOCUMENTS (INCLUDING AUTHOR, TITLE, DATE, PLACE OF PUBLICATION)						
JS	Field et al., "A novel genetic system to detect protein-protein interactions," <i>Nature</i> 340:245-246 (1989).					
JS	Gyuris et al., "Cdi1, a Human G1 and S Phase Protein Phosphatase That Associates with Cdk2," <i>Cell</i> 75:791-803 (1993).					
JS	Larsson et al., "Chromosomal Localization of Six Human Forkhead Genes, <i>freac-1</i> (FKHL5), -3 (FKHL7), -4 (FKHL8), -5 (FKHL9), -6 (FKHL10), and -8 (FKHL12)," <i>Genomics</i> 30:464-469 (1995).					
JS	Mears et al., "Autosomal Dominant Iridogoniodysgenesis Anomaly Maps to 6p25," <i>Am. J. Hum. Genet.</i> 59:1321-1327 (1996).					
JS	Mirzayans et al., "Identification of the Human Chromosomal Region Containing the Iridogoniodysgenesis Anomaly Locus by Genomic-Mismatch Scanning," <i>Am. J. Hum. Genet.</i> 61:111-119 (1997).					
JS	Nishimura et al., "Characterization of 6p25 chromosomal anomalies in two patients identifies a potential candidate gene causing congenital glaucoma," <i>Am. J. Hum. Genet.</i> 61:A21 (1997).					
JS	Nishimura et al., "The forkhead transcription factor gene <i>FKHL7</i> is responsible for glaucoma phenotypes which map to 6p25," <i>Nature Genetics</i> 19:140-147 (1998).					
JS	Orita et al., "Detection of polymorphisms of human DNA by gel electrophoresis as single-strand conformation polymorphisms," <i>Proc. Natl. Acad. Sci. USA</i> 86:2766-2770 (1989).					
JS	Sheffield et al., "Attachment of a 40-base-pair G+C-rich sequence (GC-clamp) to genomic DNA fragments by the polymerase chain reaction results in improved detection of single-base changes," <i>Proc. Natl. Acad. Sci. USA</i> 86:232-236 (1989).					
EXAMINER	J. Dumer			DATE CONSIDERED	6-21-07	
EXAMINER: Initial citation considered. Draw line through citation if not in conformance and not considered. Include copy of this form with the next communication to applicant.						



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				Applicant	Michael A. Walter et al.	
				Filing Date	April 16, 1999	
				Group	1614	
(37 CFR §1.98(b))				IDS Filed	December 13, 1999	
U.S. PATENTS						
Examiner's Initials	Patent Number	Issue Date	Patentee	Class	Subclass	Filing Date (If Appropriate)
FOREIGN PATENT OR PUBLISHED FOREIGN PATENT APPLICATION						
Examiner's Initials	Document Number	Publication Date	Country or Patent Office	Class	Subclass	Translation (Yes/No)
<i>JS</i>	WO 99/16899	04/08/99	WIPO			
OTHER DOCUMENTS (INCLUDING AUTHOR, TITLE, DATE, PLACE OF PUBLICATION)						
<i>JS</i>	Gould et al., "Autosomal Dominant Axenfeld-Rieger Anomaly Maps to 6p25," <i>American Journal of Human Genetics</i> 61(1):765-768 (1997)					
<i>JS</i>	Jordan et al., "Familial Glaucoma Iridogoniodysplasia Maps to a 6p25 Region Implicated in Primary Congenital Glaucoma and Iridogoniodysgenesis Anomaly," <i>American Journal of Human Genetics</i> 61(1):882-888 (1997)					
<i>JS</i>	Mears et al., "Mutations of the Forkhead/Winged-Helix Gene, <i>FKHL7</i> , in Patients with Axenfeld-Rieger Anomaly," <i>American Journal of Human Genetics</i> 63:1316-1328 (1998)					
<i>JS</i>	Raymond et al., "Molecular Genetics of the Glaucomas: Mapping of the First Five "GLC" Loci" <i>American Journal of Human Genetics</i> 60:272-277 (1997)					
EXAMINER	<i>S. J. J. J.</i>			DATE CONSIDERED	<i>6-21-07</i>	
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